

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

Matrix Storage and Dispensing System

FIELD OF THE INVENTION

The present invention relates to the storage and dispensing of substances. More particularly, the invention provides a system, and method of use, for serially dispensing a large number of reagents into a plurality of receptacles.

BACKGROUND OF THE INVENTION

In chemical and biological laboratories, reagent transfer from a source vessel to a target receptacle is a fundamental task. Typically, a technician must retrieve various reagent bottles from a storage location, each containing a substance pertinent to the task at hand. The technician then manually pipettes a precise quantity of each into an appropriate reaction receptacle, such as a selected well of a multi-well plate. To prevent contamination, the pipette tip must be cleaned after contact with each different reagent, or it must be discarded and replaced with a new tip.

Alternatively, the technician can attempt to manually pour each of the collected reagents from its storage vessel into a desired reaction receptacle. However, given the ultra-small quantities of reagents typically called for in modern-day protocols, particularly for expensive reagents, this technique can be very tedious and difficult to accurately perform. Moreover, the act of pouring often leads to wasted reagent, e.g., where excessive amounts are inadvertently dispensed, and cross-contamination between receptacles can result, especially when working in a high-density receptacle format (e.g., a plate or tray having ninety-six wells).

Thus, it is not surprising that such manual techniques fail to meet the demands of most laboratories, where very small quantities of numerous (e.g., hundreds or thousands) reagents must be dispensed in a quick and accurate manner.

While systems are known that automate certain aspects of reagent storage, retrieval and/or dispensing, these too are associated with certain disadvantages. One such system, available from Sagian Inc. (Indianapolis, Indiana), automates the picking and placing of reagents. Briefly, to "pick" a reagent is to retrieve it from a reagent file, and to "place" it is to re-file it back into the reagent file. The Sagian system employs two industrial robots to move reagents to and from an operator area. The first robot is a

mini-trieve that moves to a vertical file holding a target reagent and then pulls out an appropriate drawer containing the reagent. The robot then delivers the drawer to another work area where a CRS articulated robot removes the requested reagent, verifies that it is the correct container by passing the container in front of a bar-code scanner, and places it into one of a series of racks which are accessible by the operator. The mini-trieve then returns the drawer to its original location in the file. While eliminating much of the labor burden and handling errors generally associated with manual techniques, manual intervention is nevertheless required in order to dispense the reagent. Moreover, much wasted effort is involved since each drawer retrieved by the robot usually contains hundreds of additional reagents that do not pertain to the task at hand. Further, the robotic motions involved, and distances traversed, in retrieving each reagent can be quite substantial. Cumulatively, the overall process can be quite time consuming, particularly in situations where a great number of reagents (e.g., hundreds or thousands) must be retrieved.

Another automated system is sold under the trade name HAYSTACK, available from The Automation Partnership Group plc (Melbourn Science Park, Melbourn, Royston, Hertfordshire, UK). Similar to the Sagian system, the HAYSTACK system utilizes industrial robots to retrieve drawers of reagents from vertical files. In addition to such pick-and-place functions, The Automation Partnership offers modules that are able to carry out various dispensing steps. Such added capability, however, substantially increases the operational complexity of the system, and can consume a great deal of valuable laboratory space, as well.

There is, thus, a need for a relatively simple and compact reagent storage and dispensing system that provides for variable (custom) retrieval, as well as quick and accurate dispensing, of numerous selected reagents.

SUMMARY OF THE INVENTION

In one of its aspects, the present invention provides a system for storing and dispensing a plurality of reagents.

According to one embodiment, the system includes an addressable array of reagent dispensers, each having a gate mechanism disposed at a lower outlet region thereof. The gate mechanisms are independently operable between (i) an opened condition permitting passage of a respective reagent through the outlet region, and (ii) a

closed condition whereat such passage is blocked. A first support is disposed below the dispenser array, and a second support, having a holding area for receiving a plurality of receptacles, is mounted on the first support. The first and second supports can be, for example, independently operable xy stages. The first support is variably positionable in a manner permitting placement of a fixed target region thereof directly under any selected one of the dispensers in the array. The second support is variably positionable in a fashion permitting placement of any selected target site of the receptacle-holding area directly over the fixed target region.

Each of the dispensers can be, for example, an elongated container having a longitudinally extending passageway configured to receive and hold a respective reagent when the gate mechanism is in the closed condition.

A rack having an array of holding cells can support the containers. According to one embodiment, the rack has at least 100 holding cells, and preferably in excess of 1,000 holding cells. Exemplary racks include, for example, 5,000, 10,000, 50,000, 100,000 and 500,000 holding cells. Each holding cell can be configured to removably support one of the containers in a substantially upright fashion. The holding cells can be configured to hold the containers at an average density, for example, of between about 2-8 containers per cm^2 , or higher. In one embodiment, the containers are disposed in the rack at an average density of between about 3-6 containers per cm^2 ; and preferably between about 4-5 containers per cm^2 . Multiple racks (e.g., 2, 3, 4, 5, or more) can be arranged in tandem for use in an "assembly line" type fashion.

A plurality of different reagents can be disposed in the dispensers. In one embodiment, each dispenser contains a reagent that is unique to the array.

Beads can be employed to carry the different reagents. One embodiment of the invention provides a plurality of bead groups, or "lots," with each lot being comprised of substantially similar beads carrying a respective one of the different reagents. The beads can be relatively large, e.g., about 1-5mm in diameter; or the beads can be relatively small, e.g., each having a diameter of less than about a millimeter. In one preferred embodiment, each bead has a diameter of between about 275-325 μm ; and preferably about 300 μm .

In one embodiment, a plurality of reagent-carrying beads are held in sealed ampules. In an exemplary arrangement, the ampules are dimensioned to move

downward through a dispenser passageway under the force of gravity, in a substantially single-file fashion. Preferably, all of the beads in any given ampule carry the same, or a substantially similar, kind of reagent. Further in this embodiment, each passageway of the dispenser array is loaded with a plurality of such ampules.

5 One embodiment of the invention provides a detection assembly adapted to detect the passage of reagent dispensed from any one of the dispensers in the array. To this end, the detection assembly is provided with a field of view extending between the dispenser outlet regions and the second support.

According to one particular embodiment, the detection assembly includes a
10 radiation emitter, such as a diode laser, and a radiation sensor. In an exemplary arrangement, the radiation emitter is (a) mounted on the first support at a region along one side of the second support, and (b) configured to project a substantially linear radiation beam along a pathway that passes over the fixed target region of the first support. The radiation sensor can be (a) mounted on the first support at a region along
15 an opposing side of the second support, and (b) disposed within the radiation-beam pathway.

In one embodiment, each gate mechanism of the array is subject to a biasing force that normally urges it to the closed position, thereby preventing the passage of reagent through a respective outlet region. A release mechanism, adapted for positioning near
20 any one of the gate mechanisms, is operable to apply a secondary force of a magnitude and direction effective to override the normal biasing force so that the gate mechanism assumes the opened condition.

In one particular embodiment, each gate mechanism includes a magnetic pinch valve having first and second permanent magnets that are pivotally mounted in facing
25 relation at a respective outlet region. The magnets have lower, confronting north and south pole regions, respectively, that are normally urged toward one another by magnetic forces so as to pivot the magnets to the closed condition. Further in this embodiment, the release mechanism can be an electromagnet operable to generate a magnetic force having south and north pole portions disposed to attract the north and south pole lower
30 regions of the first and second pivotal magnets, respectively, so that they swing away from one another (i.e., to an open condition).

In another particular embodiment, each gate mechanism is a resiliently deflectable lever having a protrusion normally extending into a respective outlet region. Further in

this embodiment, the release mechanism is a rod adapted for reciprocal linear motion between a retracted position and an extended position. Upon movement toward the extended position, the rod can mechanically engage and deflect the lever, so that the protrusion is at least partially withdrawn from the outlet region (i.e., to an open
5 condition).

The system of the invention can further include a guide or funnel member located over the fixed target region of the first support, between the dispenser array and the second support. In a preferred embodiment, the guide member is disposed for movement with the first support to a position under any selected dispenser. The guide member is
10 configured to channel reagent dispensed from such dispenser to a selected site on the holding area of the second support.

In one particular embodiment, the guide member includes (i) an upper opening, or inlet, that is alignable with any one of the outlet regions for receiving reagent dispensed therefrom, and (ii) a lower opening, or outlet, through which dispensed reagent may
15 egress in route to the holding area. Preferably, the upper opening is larger than the lower opening. A conical portion can be provided between the upper and lower openings.

In another of its aspects, the present invention provides a reagent dispenser assembly.

According to one preferred embodiment, the reagent dispenser assembly includes a container adapted to receive a reagent and a gate mechanism located at a lower outlet region of the container. The gate mechanism is provided with first and second
20 permanent magnets pivotally mounted in facing relation at the lower outlet region. The pivotal magnets have lower, confronting north and south pole regions, respectively, that are normally urged toward one another by magnetic forces so as to swing them to a
25 closed condition whereat the egression of reagent from the container is substantially blocked.

In one embodiment, an electromagnet is disposed below the gate mechanism. In this embodiment, the electromagnet is operable to generate a magnetic force having south
30 and north pole portions disposed to attract the north and south pole lower regions of the first and second magnets, respectively, so that these regions swing away from one another to an opened condition. In this opened condition, the egression of reagent from the container is permitted.

Another embodiment provides a rack holding a plurality of the containers at respective locations defining an array. A first movable support is disposed below the rack, upon which the electromagnetic can be mounted.

A second movable support can be mounted on the first movable support, under
5 the electromagnet. In this embodiment, the second movable support is configured to receive and hold a multi-well plate for receiving reagents dispensed from the containers.

Still a further aspect of the present invention provides a method for loading a plurality of receptacles with one or more reagents.

According to one embodiment, the method includes the steps of

10 (i) placing the receptacles on a support under an addressable array of reagent dispensers;

(ii) selecting a dispenser equipped to dispense a desired reagent, and a receptacle for receiving the desired reagent;

15 (iii) simultaneously (a) positioning a fixed target region of the support at a location under the selected dispenser, and (b) positioning the selected receptacle at a location directly over the fixed target region of the support;

(iv) dispensing the desired reagent from the selected dispenser into the selected receptacle;

(v) detecting the desired reagent as it is dispensed from the selected dispenser;

20 and

(vi) repeating steps (ii) - (v) so that reagent is dispensed from at least one other dispenser into at least one other receptacle.

In one embodiment, each of the receptacles is a well of a multi-well tray.

In another embodiment, each of the dispensers is equipped to dispense an analyte-
25 specific reagent that is unique to the array.

In a further embodiment, at least 100 different analyte-specific reagents are dispensed from respective dispensers into respective receptacles. Other embodiments contemplate the dispensing of at least 500, 1,000, and 10,000, different reagents.

These and other features and advantages of the present invention will become
30 clear from the following description.

BRIEF DESCRIPTION OF THE DRAWINGS

The structure and manner of operation of the invention, together with the further objects and advantages thereof, may best be understood by reference to the following description taken in conjunction with the accompanying drawings, in which:

5 Figure 1 is a perspective view of a reagent storage and dispensing system, showing a dispenser poised for insertion into a holding cell of a support rack, according to an embodiment of the present invention.

Figure 2 is a perspective view, with portions broken away, showing additional details of the reagent storage and dispensing system of Figure 1.

10 Figure 3 is an exploded view showing still further details of the reagent storage and dispensing system of Figures 1 and 2.

Figure 4 is a partial side-sectional view of a dispenser of the invention holding a plurality of reagent-containing ampules.

15 Figures 5(A) and 5(B) are vertical and horizontal cross-sectional views, respectively, showing a magnetic pinch valve blocking the passage of reagent beads from a dispenser, in accordance with one embodiment of the present invention.

Figures 6(A) and 6(B) are vertical and horizontal cross-sectional views, respectively, showing an electromagnet inducing the magnetic pinch valve of Figures 5(A) and 5(B) to permit the passage of reagent beads.

20 Figures 7(A) and 7(B) are vertical and horizontal cross-sectional views, respectively, showing a magnetic pinch valve blocking the passage of a fluidic reagent from a dispenser, in accordance with a further embodiment of the present invention.

25 Figures 8(A) and 8(B) are vertical and horizontal cross-sectional views, respectively, showing an electromagnet inducing the magnetic pinch valve of Figures 7(A) and 7(B) to permit the passage of an aliquot of fluidic reagent.

Figure 9(A) is a side cross-sectional view showing a spring-biased lever blocking the passage of reagent-carrying ampules from a dispenser, in accordance with one embodiment of the present invention.

30 Figure 9(B) is a side cross-sectional view showing a rod-like actuator deflecting the spring-biased lever of Figure 9(A), so that a single reagent-carrying ampule can fall from the dispenser into an underlying guide or funnel member.

Figure 10 is a perspective view of the reagent storage and dispensing system of the present invention in the context of a larger system for loading microcard wells with reagent-carrying beads.

5 DETAILED DESCRIPTION OF THE INVENTION

The following discussion of the preferred embodiments of the present invention is merely exemplary in nature. Accordingly, this discussion is in no way intended to limit the scope of the invention.

One aspect of the invention provides a system for dispensing a plurality of
10 reagents. With initial reference to the embodiment of Figures 1-3, the system generally includes a movable table or platform assembly, denoted as 12, disposed under an addressable array of reagent dispensers, as at 16, equipped to serially dispense a plurality of reagents. As used herein, the wording "addressable array" refers to an array having a known reagent associated with a known location (address) in the array.

15 Platform assembly 12 includes an upper support 22 mounted on a lower support 26. Lower support 26 is movable such that a fixed (i.e., constant) target or reference region thereof, e.g., as indicated at 26a in the exploded view of Figure 3, can be positioned below any selected dispenser of array 16. Upper support 22 is movable such that any selected (i.e., variable) target site of a receptacle-holding area thereof, visible as
20 stippled region 22a in the embodiment of Figure 3, can be positioned over the fixed target region of lower support 26.

Briefly, in operation, the fixed target region of the lower support is positioned under a dispenser holding a desired reagent. At the same time, a selected target site of the upper support's receptacle-holding area is positioned over the lower support's fixed
25 target region. Typically, a particular receptacle held in a specific place on the receptacle-holding area, such as a well of multi-well plate 36, will be situated over the selected target site. Dispensed reagent, then, will fall toward the selected target site, landing in the receptacle. This procedure can be repeated to load other selected receptacles with desired reagents.

30 More particularly, the upper and lower supports, which can be xy positioners, e.g., stages, tables or similar devices, are adapted for variable positioning along respective, generally horizontal planes. Such positioning can be effected using automated means, e.g., motorized assemblies, or it can be manually effected. In one preferred

embodiment, each of two xy stages is disposed in mechanical communication with a respective computer-controlled stepper motor (not shown) via a respective screw arrangement. Suitable xy stages and controllers are available commercially, for example, from NSK Inc. of Japan.

5 A control computer (not shown) integrates the operation of the stages, for example through a program written in an event driven language such as LABVIEW® or LABWINDOWS® (National Instruments Corp., Austin, Texas). In particular, the LABVIEW software provides a high level graphical programming environment for controlling instruments. U.S. Pat. Nos. 4,901,221; 4,914,568; 5,291,587; 5,301,301;
10 5,301,336; and 5,481,741 (each expressly incorporated herein by reference) disclose various aspects of the LABVIEW graphical programming and development system. The graphical programming environment disclosed in these patents allows a user to define programs or routines by block diagrams, or "virtual instruments." As this is done, machine language instructions are automatically constructed which characterize an
15 execution procedure corresponding to the displayed procedure.

Interface cards for communicating the computer with the motor controllers are also available commercially, e.g., from National Instruments Corp.

The receptacle-holding area of the upper support is adapted to removably support a plurality of receptacles for receiving respective reagents from the dispenser array.
20 Along the holding area, means are provided for maintaining each receptacle in a desired location while the support is moved from one place to another. For example, a slightly recessed trough-like region can extend below the uppermost surface of the support, into which the receptacles can be placed. Alternatively, or in addition, mechanical holding means such as clips, brackets, bumpers, framing, VELCRO®, or the like, and/or magnetic
25 holding means, such as magnetic strips on the holding-area surface and a magnetically attractable undersurface on the receptacles, or the like, can be employed to maintain the containers in place.

In the embodiment of Figures 1-3, the receptacles are provided as an array of spaced-apart receiving wells, such as wells 32, formed in a tray or plate 36. Each of
30 wells 32 has an opening at its upper end, permitting the well to receive and hold a reagent dispensed from above. A spring-loaded plate holder (not shown), attached to the

upper surface of support 22 on opposing sides of holding area 22a, prevents plate 36 from sliding across upper support 22 as it is moved.

As previously indicated, lower support 26 is provided with a fixed (constant) reference or target region, such as area 26a visible in Figure 3. The fixed target region is a specific portion of the lower support (i) that is positionable under any dispenser of the dispenser array, and (ii) over which any selected (variable) site of the receptacle-holding area can be positioned. Typically, placement of the fixed target region will be determined by the presence of one or more elements, discussed below, each having a position and/or operational range of motion that is substantially fixed above a particular area of the lower support's upper surface. For example, the fixed target region can lie under a radiation beam, such as beam 38 in Figures 1-3, projectable over the receptacle holding area for detecting the passage of reagent from an overhead dispenser into a receptacle. As another example, the fixed target region can be located below a guide or funnel member, as at 40 in Figures 2-3, for channeling reagent dispensed from an overhead dispenser to a selected site on the receptacle holding area. As yet a further example, the fixed target region can be positioned in the vicinity of a release mechanism or actuator, as at 44 in Figures 2-3, for causing a selected reagent dispenser to dispense a desired reagent. Where more than one of the above components are employed, they will typically all be located in the general area at or above the fixed target region. Details of such components are discussed more fully below.

Turning now to the reagent dispenser array, each reagent dispenser takes the form of an elongated container, such as cylindrical or tubular container 42 shown poised above array 16 in Figure 1. The containers can be formed, for example, of plastic, glass, and/or metal, or other material. In one embodiment, each container is a rigid cylinder formed of a metal or metal alloy (e.g., aluminum, an aluminum alloy, or stainless steel), intended for repeated uses. In another embodiment, each container is constructed of a relatively inexpensive material, such as glass or plastic that can be readily disposed of after its contents (reagent) have been exhausted.

By configuring each container with a sufficiently narrow diameter, a high density of such containers can be achieved. For example, various embodiments contemplate from about 2 to 8 containers per cm^2 , on average, or higher. One preferred embodiment contemplates an average density of between about 3-6 containers per cm^2 ; and most preferably between about 4-5 containers per cm^2 . In an exemplary arrangement, a

plurality of substantially like containers, each having a diameter of less than 1cm, are disposed with substantially parallel longitudinal axes and at closely spaced positions defining an array. In one particularly preferred embodiment, an array of such containers, each having an outer diameter of about 4mm, are arranged with a center-to-center spacing between adjacent containers of about 4.50mm.

Each container is provided with a passageway configured to receive and hold a respective reagent. In the embodiment of Figure 1, a longitudinally extending lumen, denoted as 42a, holds a plurality of reagent-containing ampules, such as 50. The passageway can be of any horizontal cross-section, such as circular, oval, polygonal, or other cross-section. Optionally, the exposed inner sidewalls of the passageways can be covered with a substantially inert lining material.

A rack or frame, generally denoted as 46, provides a plurality of holding cells, each being configured to support one reagent container therein. In Figure 1, for example, container 42 can be inserted into one of holding cells 52 of frame 46 by lowering it in the direction of the darkened arrow. The rack can have any number of holding cells. In one embodiment, the number of different reagents held in the rack determines the number of holding cells. That is, there can be a one-to-one correspondence between the number of holding cells and the number of different reagents. For situations requiring a relatively large quantity of a particular reagent, other embodiments provide such reagent in two or more holding cells of the array.

Rack 46 can have tens, hundreds, thousands, tens of thousands, or hundreds of thousands of holding cells. Advantageously, such configurations permit the storage and variable selection of many different reagents. In one particularly preferred embodiment, rack is formed with 10,000 holding cells, each removably supporting a respective reagent container in a substantially upright fashion. In this embodiment, any combination of up to 10,000 different reagents can be dispensed into the wells of a multi-well tray.

It should further be appreciated that a plurality of such racks can be utilized, for example, in an "assembly line" type arrangement. For example, three 10,000 cell racks can be arranged at respective locations along an automated system, each capable of dispensing up to 10,000 different reagents.

The particular form of each reagent stored and dispensed in accordance with the teachings of the invention is not critical, provided only that it is compatible with the storage and dispensing means. The reagent, which can be a single substance or a

grouping of different substances, can be provided, for example, as a solid, liquid, powder, emulsion, suspension or substantially any combination thereof. In one embodiment, a coating material is applied to a reagent core to form particulates, pills, beads or tablets. The coating can be dissolvable or swellable to permit access to the reagent under controllable conditions (e.g., upon exposure to a particular solvent).

Guidance for preparing coated micro-particles (beads) is provided, for example, in: [1] R. Pommersheim, H. Lowe, V. Hessel, W. Ehrfeld (1998), "Immobilization of living cells and enzymes by encapsulation," Institut für Mikrotechnik Mainz GmbH, IBC Global Conferences Limited; [2] F. Lim A. Sun (1980), *Science* 210, 908; [3] R. Pommersheim, J Schrezenmeir, W. Vogt (1994), "Immobilization of enzymes and living cells by multilayer microcapsules" *Macromol Chem. Phys* 195, 1557-1567; and [4] W. Ehrfeld, V. Hessel, H. Lehr, "Microreactors for Chemical Synthesis and Biotechnology-Current Developments and Future Applications" in: *Topics in Current Chemistry* 194, A. Manz, H. Becker, *Microsystem Technology in Chemistry and Life Science*, Springer Verlag, Berlin Heidelberg (1998), 233-252; each expressly incorporated herein by reference.

In another embodiment, a plurality of bead-like particles act as solid supports for the reagents. For example, reagents can be synthesized on the beads, or absorbed thereto. In still a further embodiment, a slurry or dispersion comprised of a reagent and binding material is used to form a plurality of bead-like particles, with each individual bead having a substantially homogenous consistency.

A plurality of different reagents can be formed into respective collections or groups of reagent beads, or "lots." For example, 10,000 different reagents can be formed into 10,000 different bead lots, with each lot comprised of a plurality of substantially like beads carrying a respective reagent. Beads from each lot can then be loaded into respective dispensers of the dispenser array.

In one embodiment, a plurality of bead lots are formed, wherein each bead includes a reagent core covered with a coating material, such as a gelatin, having well-defined physical and chemical properties. Preferably, in this embodiment, all beads in all lots bear substantially the same outer coating (i.e., a "generic" coating). It should be appreciated that this arrangement reduces the risk of equipment contamination due to contact with the reagents. If any residues are left behind as the reagents move through

the system, such residues will all be of the same coating material. Preferably, the coating material is chosen so that any residues are innocuous to the system.

Further regarding reagent-carrying beads, the beads can be formed with a diameter slightly less than that of one of the passageways of the dispenser array, so that the beads can be stacked in each container, one on top of the other, for gravity-fed dispensing. For example, beads having a diameter of between about 3.50-3.90mm, and preferably about 3.70mm, can be stacked in a container having a passageway with a diameter of about 4mm.

Alternatively, the beads can be relatively small, e.g., each having a diameter of less than about 1mm. In one preferred embodiment, each bead has a diameter of between about 275-325 μ m, and preferably about 300 μ m. A plurality of such beads can be placed in a capsule or ampule to be dispensed as a unit. For example, hundreds or thousands of beads from the same or substantially identical lots can be packed into plastic ampules. The exemplary arrangement of Figure 4 shows reagent-carrying beads 62 disposed in bullet-shaped ampules 50. The various passageways in the dispenser array can be loaded with a plurality of such ampules, each containing beads from respective lots. As best seen in Figure 4, the ampules can be dimensioned to move downward through the passageways under the force of gravity in a substantially single-file fashion.

The ampules can be provided with a cover member over an upper opening thereof. The cover member can be, for example, a removable cap or dome having an open end configured to fit snugly about the upper region of an ampule. Or, a frangible sheet-like film or membrane, such as membranes 66 in Figure 4, can be applied to an upper rim or lip surrounding the upper opening of each ampule. Access to the beads can be gained, for example, by removing or rupturing the membrane cover.

In one embodiment, the cover over each ampule forms a substantially airtight seal, protecting the reagent contents of the ampule from the ambient atmosphere. The seal can be effected, for example, using conventional adhesives or by heating-sealing techniques. The sealed ampules can further contain an inert gas, such as nitrogen or the like.

Substantially any reagent can be stored and dispensed using the system of the present invention. According to one preferred embodiment, the reagent in each dispenser includes components useful for real time fluorescence-based measurements of nucleic acid amplification products (such as PCR) as described, for example, in PCT Publication

WO 95/30139 and U.S. Patent Application Serial No. 08/235,411, each of which is expressly incorporated herein by reference.

In one embodiment, each container holds an analyte-specific reagent effective to react with a selected analyte that may be present in a sample. For example, for
5 polynucleotide analytes, the analyte-specific reagent can include first and second oligonucleotide primers having sequences effective to hybridize to opposite end regions of complementary strands of a selected polynucleotide analyte segment, for amplifying the segment by primer-initiated polymerase chain reaction. The analyte-specific
10 detection reagent can further include a fluorescer-quencher oligonucleotide capable of hybridizing to the analyte segment in a region downstream of one of the primers, for producing a detectable fluorescent signal when the analyte is present in the sample.

An accession or tracking number can be printed on each container, identifying the reagent contained therein. For those embodiments employing ampules to hold the reagents, each ampule can bear such a tracking number. With regard to the latter, the
15 containers can be formed with window regions through which the tracking numbers on the ampules can be observed. The window regions can be of a transparent material, such as glass or plastic, or they can be openings or notches formed in the sidewalls of the containers.

Preferably, each tracking number is provided in a machine-readable format, such
20 as a bar code. An operator can manually scan the bar codes, or they can be scanned in an automated fashion using robots. In one embodiment, a robot picks up a container from a tray of reagent tubes and wands a bar code to learn and/or confirm the identity of the reagent held therein. Using the scanned information, a control computer instructs the robot to place the tube in a designated holding cell of a rack. Alternatively, or in
25 addition, the bar codes on the ampules can be scanned using a robot as they are dispensed from the containers, and the bar codes on the containers can be manually scanned as they are inserted into or removed from the holding cells in rack 46.

Controllable dispensing of each reagent is provided by a gate mechanism located at a lower outlet region of each dispenser. Each gate mechanism is independently
30 operable between (i) an opened condition permitting passage of a respective reagent through the outlet region, and (ii) a closed condition whereat such passage is blocked. The particular construction of the gate mechanism is not critical, provided only that it is capable of retaining the reagent held in the respective container until such time that it is

desired to dispense the reagent. Additionally, each gate mechanism is preferably operable on an individual basis, so that the various reagents can be dispensed one at a time.

Several exemplary gate mechanisms for use in connection with various types of reagents will now be described.

According to one embodiment, each gate mechanism includes a magnetic pinch valve having first and second magnets that are pivotally mounted in facing relation at a respective container outlet region. Generally, the pinch valve magnets have lower, confronting north and south pole regions that are urged toward one another by magnetic forces, thereby normally disposing the gate mechanism in a "closed" condition. Additional structure may be included to supplement or enhance such normal positioning of the pivotal magnets.

Figures 5 and 6 show one particular embodiment of a magnetic pinch valve that is especially useful for dispensing reagent-carrying beads. Here, a supportive insert or plug, such as 170, is disposed in a lower region of each holding cell 152 of rack 146. Frictional engagement of the insert's outer sidewall with the inner sidewall of a respective holding cell can hold the insert in place. Adhesives or other retaining means may be employed to ensure the long-term placement of each insert. The lower end of an elongated container 142, containing reagent beads 188, rests on an upper, inwardly flanged portion of insert 170b. Insert 170 provides pivot points, denoted as 174 and 176, on opposing inner sidewall portions to which respective permanent magnets 178, 180 pivotally attach at their midregions, e.g., by way of pivot pins. As best seen in Figures 5(B) and 6(B), each pivotal magnet 178, 180 is substantially C-shaped in horizontal cross-section. Pivotal magnets 178, 180 are oriented such that their upper and lower end regions are of opposite polarity. A third permanent magnet, denoted as 184, is fixedly positioned along a sidewall portion of insert 170, above pivot points 174, 176. One end of this stationary magnet 184 is disposed adjacent an upper region of one of the pivotal magnets, 178 or 180; and the other end of magnet 184 is disposed adjacent an upper region of the other pivotal magnet. Stationary magnet 184 is oriented such that the polarity of each such end is opposite that of the upper region of the pivotal magnet adjacent thereto. Accordingly, in the normal state, the upper region of each pivotal magnet 178, 180 is attracted to an adjacent portion of the sidewall-mounted magnet 184 and, at the same time, the confronting north and south pole regions of the pivotal

magnets are attracted toward one another. In response, magnets 178, 180 pivot about their respective pivot points 174, 176 so that their lower north and south pole regions swing toward one another; thereby assuming the closed condition, as shown in Figures 5(A) and 5(B). In the closed condition, the outlet region is constricted such that the reagent beads 188 are not able to fall out.

To release a reagent bead, a release mechanism is moved to a position under the outlet region of a selected container. The release mechanism is operable to overcome the closing force that normally prevents the egression of reagent beads. In the embodiment of Figures 5-6, an electromagnet 192 is used as the release mechanism. Electromagnet 192 has spaced-apart south and north pole portions disposed to attract the opposing north and south pole lower regions of pivotal magnets 178, 180, respectively, in a direction away from one another. The magnetic force generated upon activating electromagnet 192 is sufficient to overcome the previously-described normal closing force, thereby swinging the lower regions of pivotal magnets 178, 180 apart so that one of the beads 188 can fall through a central opening 170a at the bottom of insert 170. As best seen in Figure 6(A), upon swinging the lower regions of pivotal magnets 178, 180 apart, the upper regions swing toward one another, thereby blocking the passage of any additional beads 188. Once a bead has been dispensed, electromagnet 192 can be deactivated, permitting the gate mechanism to return to the closed position, as shown in Figures 5(A)-5(B). Electromagnet 192 can then be moved to another container for dispensing another reagent. It should be appreciated that this arrangement allows for the realization of controllable, single-bead dispensing.

Figures 7 and 8 show an embodiment of a pinch valve especially useful for dispensing a fluidic reagent. In this embodiment, two substantially planar, permanent magnets 278, 280 attach at their uppermost ends to a support member 270, for swinging motion about respective pivotal connections 274, 276. As best seen in Figures 7(A) and 8(A), the pivotal connections 274, 276 are disposed on opposing sides of a lowermost opening of elongated container 242. Support member 270, in turn, is fixed to a lower end region of container 242. In this regard, an annular cavity 287 extends upwardly from a lowermost rim or lip of container 242, circumscribing longitudinal passageway 242a. An upstanding cylindrical collar 270a, formed at the top of support member 270, is configured to fit snugly into cavity 287. Collar 270a can be maintained in cavity 287 by frictional forces and/or adhesive agents. A plate 272 is secured against a lowermost

end of holding cell 252 to provide a lower foundation for supporting the container and gate assembly therein.

Gasket members 279, 281 (Figures 7(B) and 8(B)) are affixed to opposite inner sidewall portions of support member 270. Gasket members 279, 281 provide opposing planar surfaces positioned for sliding, substantially fluid tight, contact with the lateral side-edges of pivotal magnets 278, 280. The opposing planar surfaces of gasket members 279, 281 can be formed of a hydrophobic material, and/or treated to exhibit hydrophobic characteristics, to discourage undesired leakage of the liquid reagent 288.

Similar to the previous embodiment, pivotal magnets 278, 280 are oriented such that they having lower end regions of opposite polarity. So arranged, the lower ends of pivotal magnets 278, 280 are normally attracted such that they swing toward one another and make contact, establishing a substantially fluid-tight seal (i.e., a "closed" position). In this regard, one or both magnets 278, 280 can bear a polymeric coating (not shown) along the region of contact to assist in the formation of the fluid-tight seal.

To release the liquid reagent, a release mechanism is moved to a position under the outlet region of a selected container. With reference now to Figures 8(A)-8(B), an electromagnet 292 is employed as the release mechanism. Electromagnet 292 has spaced-apart south and north pole portions disposed to attract the opposing north and south pole lower regions of pivotal magnets 278, 280, respectively, in a direction away from one another. The magnetic force generated upon activating electromagnet 292 is sufficient to overcome the previously-described normal closing force, thereby swinging the lower regions of pivotal magnets 278, 280 apart so that an aliquot of fluidic reagent 288 can fall through a central opening 272a in plate 272. The duration of activation of electromagnet 292 can be used to gauge the amount of liquid reagent dispensed. When electromagnet 292 is turned off, the normal attraction between the opposing lower end regions of magnets 278, 280 returns the valve to a closed position.

In another embodiment, each gate mechanism is a resiliently deflectable lever having a protrusion that normally extends into a respective outlet region. In an exemplary arrangement, as illustrated in Figures 9(A)-9(B), an elongated lever, indicated generally by the reference numeral 302, extends longitudinally along the outer sidewall of container 342, proximate the container's lower outlet region. Upper and lower nubs or protrusions, denoted respectively as 302b and 302c, project outwardly from one side of lever 302, towards container 342. Container 342, in turn, is provided with upper and

lower bores, indicated respectively as 342b and 342c, that extend fully through its sidewall at locations adjacent to lever 302. More particularly, each of bores 342b, 342c is positioned in alignment with a respective one of protrusions 302b, 302c, and is configured to removably receive such protrusion therein.

5 A resilient spring member, such as leaf spring 351, is secured at one of its ends to a support structure 353 that is fixedly positioned near the lower outlet region of container 342. The other end of leaf spring 351 is disposed to act against a side of lever 302, opposite container 342. Leaf spring 351 provides a normal biasing force, along the direction "F," that presses the lower end region of lever 302 against container 342.

10 Under these circumstances, lower protrusion 302c extends through lower bore 342c and into passageway 342a at the container's lower outlet region, as shown in Figure 9(A). In this "closed" position, lower protrusion 302c blocks the egression of any reagent-carrying ampules 388a-388c. Upper protrusion 302b, on the other hand, is positioned outside of passageway 342a in the normal, closed position.

15 To release an ampule, a release mechanism is moved to a position under the outlet region of a selected container. The release mechanism is operable to overcome the closing force that normally prevents the egression of ampules. With reference to Figures 9(A) and 9(B), an elongated rod 372 can be employed as the release mechanism. Rod 372 is adapted for reciprocal linear motion between (i) a retracted position, whereat rod 372 is positionable below a selected lever, such as lever 302 in Figure 9(A), to (ii) an
20 extended position, whereat a rounded-head portion of rod, denoted at 372a, can abut and press against a lower, outwardly-angled cam surface of lever, as at 302a in Figure 9(B), thereby deflecting lever 302 away from container 342. Such motion of rod 372 can be effected, for example, by providing rod at the end region of a movable plunger extending
25 from a conventional solenoid assembly.

Notably, when lever 302 is deflected in the manner just described, lower protrusion 302c is withdrawn from passageway 342a, permitting the bottommost reagent-carrying ampule 388a to fall from the container's lower outlet region. Also during such deflection, upper protrusion 302b is received within upper bore 342b such that it extends
30 into passageway 342a, preventing the egression of any remaining ampules 388b-388c. Upon returning rod 372 to its retracted position, lever 302 reassumes its normally closed position, as in Figure 9(A), preventing the passage of any ampules. It should be appreciated that this arrangement permits controllable, single-ampule dispensing.

In an alternative embodiment, similar to the embodiment just described, the release mechanism operates according to magnetic principles. In an exemplary arrangement (not shown), the lower end of the resiliently deflectable lever and the upper head portion of reciprocally movable rod are magnetically polarized, or polarizable, to exhibit the same polarity (e.g., both being "N"). The lever can be deflected by moving the rod into proximity with the lever's lower end, so that the like magnetic pole portions repel one another. Notably, contact between the rod and the lever is not required in this embodiment. The magnetic repulsion is sufficient to deflect the lever away from the container, thereby permitting a reagent-carrying ampule to fall from the container's lower outlet region.

Any of the above-described release mechanisms can be adapted for variable positioning along a generally horizontal plane under the dispenser array by mounting it to the upper surface of the lower xy stage. In this regard, a mounting assembly, such as bracket 51 of Figures 1-3, can be used to hold the release mechanism. In this embodiment, one end of bracket 51 is affixed to the upper surface of lower xy stage 26. The other (free) end of bracket 51 extends into the region between the platform assembly 12 and the dispenser array 16, whereat the release mechanism is supported, as schematically indicated at 44.

A detection assembly can be provided for detecting the passage of reagent into a receptacle from a selected overhead dispenser. In the exemplary arrangement of Figures 1-3, the region between a radiation source or emitter, such as laser 37, and a radiation sensor, as at 39, defines a detection zone. In this embodiment, both laser 37 and sensor 39 are fixedly positioned at respective locations on the upper surface of the lower xy stage 26, on opposing sides of upper xy stage 22. This construction permits movement of the detection zone along a generally horizontal plane under dispenser array 16 with movement of the lower xy stage 26. Thus, by moving xy stage 26 in an appropriate manner, the detection zone can be placed under any selected dispenser of array 16. A narrow-width beam, indicated at 38, can be directed from laser 37 along the detection zone and detected by sensor 39. An interruption in the beam 38 indicates the passage of a reagent from a dispenser above the detection zone.

Lasers and sensors, suitable for use in practicing the invention, are available commercially, for example, from Edmund Scientific (Barrington, NJ). A particularly

preferred diode laser, for use as a radiation emitter, is made by Coherent, Inc. (Auburn, CA).

In addition to detecting the passage of reagent, beam 38 can also be employed to confirm the identity of a dispensed reagent. For example, beam 38 can "double" as a bar-code scanner. In one embodiment, each reagent-carrying ampule bears a bar code that is unique to the particular type of reagent held therein. As a dispensed ampule passes through the beam, the bar code is read and the information is passed on to the control computer. The computer can then positively identify the dispensed reagent, and take appropriate corrective measures in the event of a dispensing error.

A guide or funnel member can be provided in the region between the dispenser array and the platform assembly for channeling reagent dispensed from an overhead dispenser to a selected site on the receptacle-holding area of the upper xy stage. In the exemplary arrangement of Figures 1-3, such a guide member, indicated schematically at 40, is fixedly positioned relative to the upper surface of lower support by way of mounting bracket 51. This construction permits movement of the guide member 40 along a generally horizontal plane under the dispenser array 16 with movement of the lower xy stage 26.

Typically, in operation, guide member 40 will be positioned under a selected dispenser of array 16. A selected receptacle, such as a particular well of multi-well plate 36, will be positioned under the guide member by appropriately moving the upper xy stage 22. Such positioning of the guide member and the receptacle will preferably occur simultaneously. A dispensed reagent, then, will fall through a central, vertically extending channel of guide member 40 on its way from the selected dispenser to the selected receptacle.

In one particular embodiment, shown in Figures 9(A) and 9(B), a guide member, denoted as 340, includes (i) an upper opening 340a, (ii) a lower opening 340c, smaller than the upper opening 340a, and (iii) a conical or funnel-shaped portion 340b, between the upper and lower openings. Also in this embodiment, it should be noted that the guide member 340 and reciprocally-movable rod 372 are conveniently provided together in a common housing.

At this point, the significance of the fixed target region of the lower support can be well appreciated. The fixed target region is primarily a reference point that, when positioned under a selected dispenser, facilitates the proper and simultaneous positioning

of one or more additional components thereunder. For example, with reference to the embodiment of Figures 1-3, each of the following elements is positioned at an appropriate location with respect to a selected container, as described, upon moving the fixed target region 26a to a location under such container:

5 (i) radiation beam 38 is located under the selected container so that any reagent dispensed from the container will break the beam;

(ii) guide or funnel member 40 is disposed with its upper, large opening axially aligned with the lower outlet region of the selected container so that dispensed reagent will fall therein; and

10 (iii) release mechanism 44 is positioned proximate a normally closed gate mechanism at the lower outlet region of the selected container.

Moreover, a selected well of multi-well plate 36, supported at the receptacle-holding area of the upper xy stage 22, can be moved to a position over the fixed target region while the fixed target region is being moved under the selected receptacle as just
15 described. In this way, the receptacle, too, can be quickly and accurately positioned to receive a dispensed reagent.

As with the xy stages 22, 26, operation of the various components and sub-assemblies described above can be controlled and orchestrated using the LABVIEW® or LABWINDOWS® software from National Instruments (Austin, Texas) by techniques
20 known in the art.

In a typical use, an array of holding cells 52 in rack 46 are loaded with respective containers 42, each holding a particular reagent. A data set or table is created comprised of values identifying each location or address of the holding-cell matrix with its particular resident reagent. The data set is stored electronically on a drive unit accessible to a
25 control computer. The reagents are maintained in rack 46 until dispensed, e.g., as follows.

Multi-well plate 36 is placed on receptacle-holding area 22a of upper xy stage 22. A plurality of reagents, stored in rack 46, are selected for dispensing into chosen wells 32 of plate 36, and this information is fed to the control computer. The computer
30 accesses the data set of location information to determine which containers hold the selected reagents, and a loading sequence is constructed and held in memory. The computer signals the motor controller to move lower xy stage 26 to a location whereat its

fixed target region 26a is disposed under the first container of the loading sequence. At the same time, upper xy stage 22 positions a selected target region of the receptacle-holding area 22a, underlying a chosen well of multi-plate 36, over the fixed target region 26a. Together, these steps serve to position radiation beam 38, guide member 40, and release mechanism 44 at appropriate respective locations proximate a lower outlet region of the container, as well as to position the chosen well of plate 36 under the container. The computer then signals activation of release mechanism 44, causing the gate mechanism at the lower outlet region of the container to open and dispense an aliquot of its respective reagent into the chosen well of plate 36. A correctly dispensed substance will briefly prevent radiation beam 38 from reaching sensor 39, indicating successful dispensing. If radiation beam 38 is not interrupted, as expected, an error in dispensing is recorded and a further attempt at dispensing the reagent can then be made, as desired. After the first reagent has been dispensed, the next reagent of the loading sequence can be dispensed in a like manner.

In one preferred embodiment, the dispensing system of the present invention is utilized as a sub-assembly in a larger system for loading an array of very small wells in a microcard with respective reagents. In the exemplary arrangement of Figure 10, a 384-well plate, denoted as 436, serves as a source of reagent for serially loading target 384-well microcards, such as 441, supported for movement on a carousel assembly 449. The wells of both the plate and the microcards are disposed in 16 X 24 arrays, spaced about 4.5mm center-to-center. The wells of the plate, however, have a greater diameter than the wells of the microcards. For example, each well of the plate can be configured with a diameter of about 3mm, while each well of the microcards can be formed with a diameter of about 1mm.

Generally, each well of the multi-well plate 436 is loaded with a respective reagent-carrying ampule, in accordance with the foregoing detailed description. A robot, such as 443, then moves the ampule-loaded plate, in the direction of darkened arrow "P," to a pick-and-place unit 445. Pick-and-place unit 445 simultaneously retrieves a reagent bead from each ampule in the plate, and retains the beads at spaced-apart locations defining an array corresponding to the plate and microcard arrays. Pick-and-place unit 445 then rotates about its central axis to position the retained beads over the wells of a microcard supported on the carousel at a location directly adjacent thereto, at which point it releases the beads. A bead distributor, such as 447, interposed between pick-and-place

unit 445 and the target microcard, separately channels each released bead into its designated well. The loaded microcard wells can then be used to carry out a desired assay or reaction, such as real-time PCR.

Additional details of the micro-card loading system are provided in U.S.

5 application serial no. 09/250,697, filed February 16, 1999 and U.S. application, Attorney Docket No. 0550-0076.30, filed on even date herewith, the contents of which are expressly incorporated herein by reference.

The many benefits offered by the storage and dispensing system of the present invention can now be appreciated. For example, simultaneous movement of the upper
10 and lower supports (e.g., xy stages) allows quick and accurate positioning of the various system components. Moreover, once a desired receiving receptacle is positioned under a selected dispenser, the release mechanism and detection beam can operate immediately to dispense the substance. Consequently, serial dispensing of a plurality of substances can be accomplished in a very rapid manner.

15 Those skilled in the art can now appreciate from the foregoing description that the broad teachings of the present invention can be implemented in a variety of forms. Therefore, while this invention has been described in connection with particular embodiments and examples thereof, the true scope of the invention should not be so limited. Various changes and modification may be made without departing from the
20 scope of the invention, as defined by the appended claims.

It is claimed:

1. A system for storing and dispensing a plurality of reagents, comprising:
an addressable array of reagent dispensers;
5 a gate mechanism at a lower outlet region of each dispenser, each gate mechanism being independently operable between (i) an opened condition permitting passage of a respective reagent through said outlet region, and (ii) a closed condition whereat such passage is blocked;
a first support disposed below said array;
10 a second support mounted on said first support, said second support having a holding area for receiving a plurality of receptacles;
wherein (i) said first support is variably positionable in a fashion permitting placement of a fixed target region thereof directly under any selected one of said dispensers in said array, and (ii) said second support is variably positionable in a fashion
15 permitting placement of any selected target site of said holding area directly over said fixed target region.
2. The system of claim 1, wherein said dispensers are elongated containers, each having a longitudinally extending passageway configured to receive and hold a respective
20 reagent when said gate mechanism is in the closed condition.
3. The system of claim 2, further comprising a rack having an array of at least about 100 holding cells, each holding cell being configured to removably support one of said containers in a substantially upright fashion.
25
4. The system of claim 3, wherein said holding cells are disposed at an average density of at least about 3 holding cells per cm^2 .
5. The system of claim 4, wherein said array includes at least about 500 holding
30 cells; and wherein said holding cells are disposed at an average density of at least about 4 holding cells per cm^2 .

6. The system of claim 5, wherein said array includes at least about 1,000 holding cells.

7. The system of claim 6, wherein said array includes at least about 10,000 holding cells.

8. The system of claim 7, wherein said array includes at least about 100,000 holding cells.

9. The system of claim 2, further comprising a plurality of different reagents disposed in said dispenser passageways.

10. The system of claim 9, wherein each of said passageways contains a reagent that is unique to said array.

11. The system of claim 9, further comprising a plurality of bead lots, each lot comprised of a plurality of substantially similar beads carrying a respective one of said different reagents.

12. The system of claim 11, wherein each bead has a diameter of less than about a millimeter.

13. The system of claim 12, further comprising a plurality of sealed ampules, said ampules being dimensioned to move downward through any one of said passageways under the force of gravity in a substantially single-file fashion;

wherein each ampule contains beads from the same or substantially identical lots, and wherein each passageway is loaded with a plurality of such ampules.

14. The system in accordance with any one of the preceding claims, wherein said first and second supports are independently operable xy stages.

15. The system in accordance with any one of the preceding claims, further comprising a detection assembly having a field of view extending between said dispenser outlet regions and said second support, and adapted to detect the passage of reagent dispensed from any one of said dispensers.

5

16. The system of claim 15, wherein said detection assembly includes

(i) a radiation emitter that is (a) mounted on said first support at a region along one side of said second support, and (b) configured to project a substantially linear radiation beam along a pathway that passes over said fixed target region of said first support; and

10

(ii) a radiation sensor that is (a) mounted on said first support at a region along an opposing side of said second support, and (b) disposed within said radiation-beam pathway.

15

17. The system in accordance with any one of the preceding claims, wherein each gate mechanism is subject to a normal biasing force that urges it to the closed position, thereby preventing the passage of reagent through a respective outlet region.

18. The system of claim 17, further comprising a release mechanism positionable near any one of said gate mechanisms and operable to apply a secondary force of a magnitude and direction effective to override the normal biasing force so that the gate mechanism assumes the opened condition.

20

19. The system of claim 18,

25

wherein each gate mechanism includes a magnetic pinch valve having first and second magnets (i) that are pivotally mounted in facing relation at a respective outlet region and (ii) that have lower, confronting north and south pole regions, respectively, that are normally urged toward one another by magnetic forces so as to pivot said magnets to the closed condition.

30

20. The system of claim 19,

wherein said release mechanism is an electromagnet positioned adjacent said radiation-beam pathway, said electromagnet operable to generate said secondary force as

a magnetic force having south and north pole portions disposed to attract the north and south pole lower regions of said first and second magnets, respectively, so that said lower regions pivot away from one another.

5 21. The system of claim 18,
 wherein each gate mechanism is a resiliently deflectable lever having a protrusion normally extending into a respective outlet region.

10 22. The system of claim 21,
 wherein said release mechanism is a rod adapted for motion from a retracted position to an extended position adjacent said radiation-beam pathway, said rod adapted to apply said secondary force as a mechanical force by engaging and deflecting said lever, when moved to said extended position, so that said protrusion is at least partially withdrawn from said outlet region.

15 23. The system in accordance with any one of the preceding claims, further comprising

 a guide member located over said fixed target region of said first support, between said dispenser array and said second support;

20 said guide member (i) disposed for movement with said first support to a position under any selected dispenser, and (ii) configured to channel reagent dispensed from such dispenser to a selected site on said holding area of said second support.

25 24. The system of claim 23, wherein said guide member includes
 (i) an upper opening alignable with any one of said outlet regions for receiving reagent dispensed therefrom;
 (ii) a lower opening, smaller than said upper opening, through which dispensed reagent may egress in route to said holding area; and
 (iii) a conical portion between said upper and lower openings.

30 25. A reagent dispenser assembly, comprising:
 a container adapted to receive a reagent;
 a gate mechanism located at a lower outlet region of said container;

said gate mechanism including first and second permanent magnets (i) that are pivotally mounted in facing relation at said lower outlet region and (ii) that have lower, confronting north and south pole regions, respectively, that are normally urged toward one another by a magnetic force so as to pivot said magnets to a closed condition whereat the egression of reagent from said container is substantially blocked.

26. The reagent dispenser of claim 25, further comprising:

an electromagnet disposed below said gate mechanism,

said electromagnet being operable to generate a magnetic force having south and north pole portions disposed to attract the north and south lower regions of said first and second magnets, respectively, so that said lower regions pivot away from one another to an opened condition, permitting the egression of reagent from said container.

27. The reagent dispenser of claim 26, further comprising

a rack holding a plurality of said containers at respective locations defining an array; and

a first movable support disposed below said rack;

wherein said electromagnetic is mounted on said movable support.

28. The reagent dispenser of claim 27, further comprising

a second movable support mounted on said first movable support, under said electromagnet;

said second movable support configured to receive and hold a multi-well plate for receiving reagents from said containers.

29. A method for loading a plurality of receptacles with one or more reagents, comprising:

(i) placing a plurality of receptacles on a support under an addressable array of reagent dispensers;

(ii) selecting a dispenser equipped to dispense a desired reagent, and a receptacle for receiving the desired reagent;

(iii) simultaneously (a) positioning a fixed target region of the support at a location under the selected dispenser, and (b) positioning the selected receptacle at a location directly over the fixed target region of the support;

(iv) dispensing the desired reagent from the selected dispenser into the selected
5 receptacle;

(v) detecting the desired reagent as it is dispensed from the selected dispenser;
and

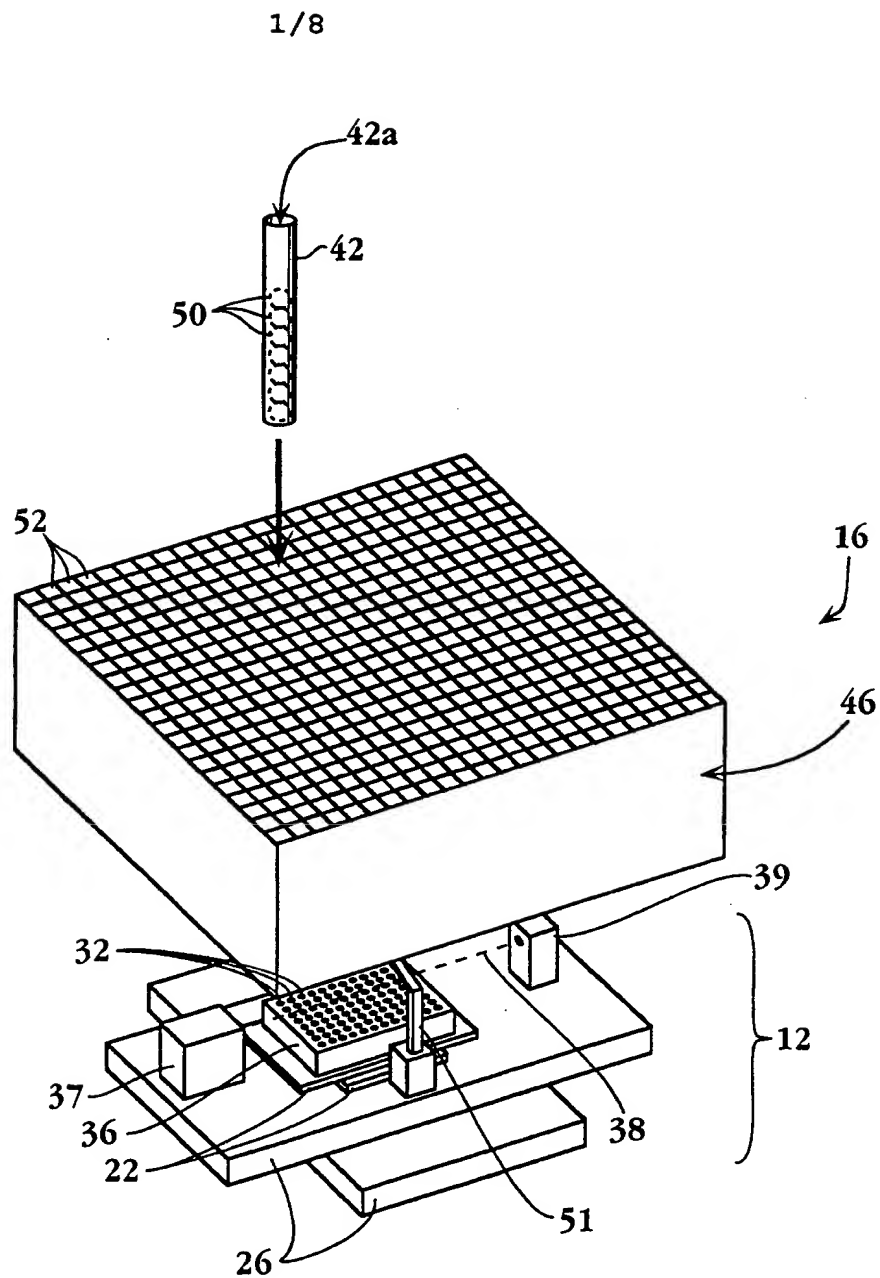
(vi) repeating steps (ii) - (v) so that reagent is dispensed from at least one other
dispenser into at least one other receptacle.

10 30. The method of claim 29, wherein each of said receptacles is a well of a multi-well tray.

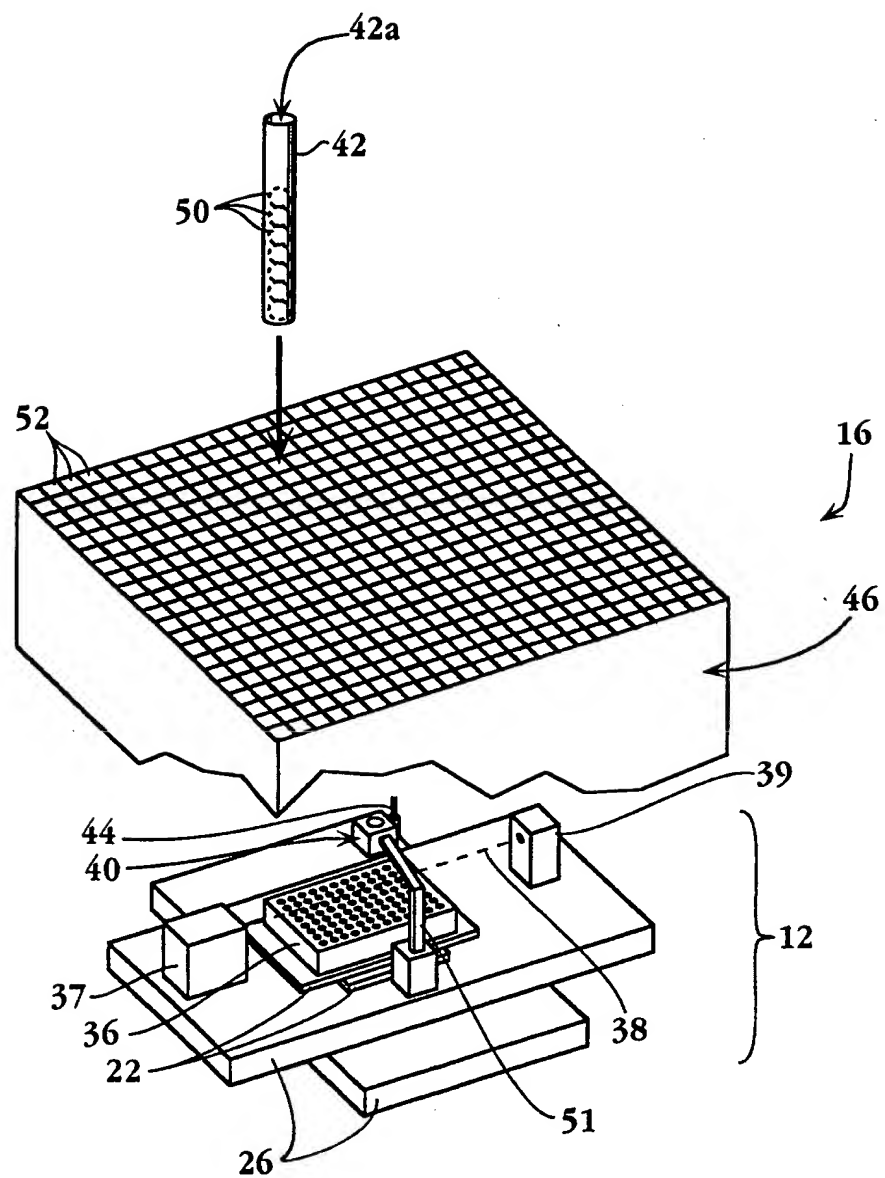
31. The method of claim 29, wherein each of said dispensers is equipped to
15 dispense an analyte-specific reagent that is unique to said array.

32. The method of claim 29, wherein at least 100 different analyte-specific reagents are dispensed from respective dispensers into respective receptacles.

20 33. The method of claim 32, wherein at least 500 different analyte-specific reagents are dispensed from respective dispensers into respective receptacles.

**Fig. 1**

2/8

**Fig 2**

3/8

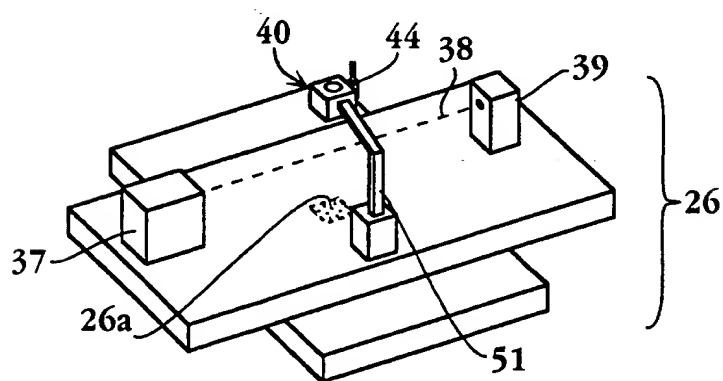
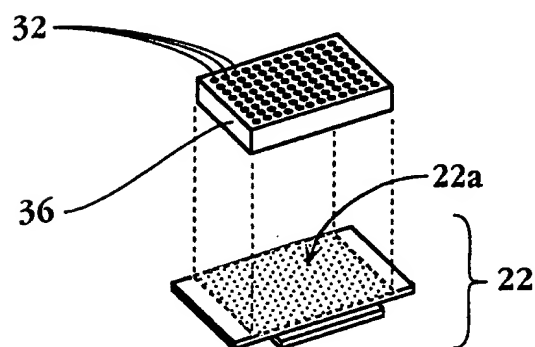
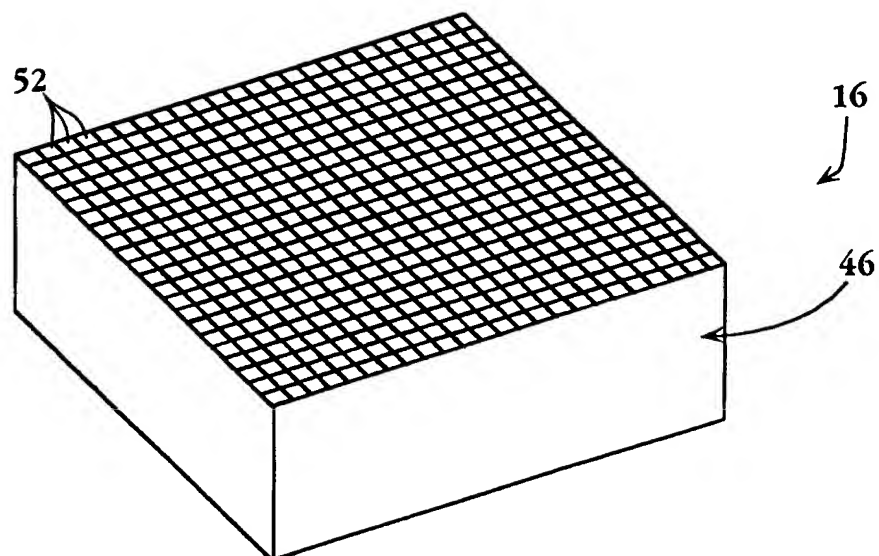


Fig. 3

4 / 8

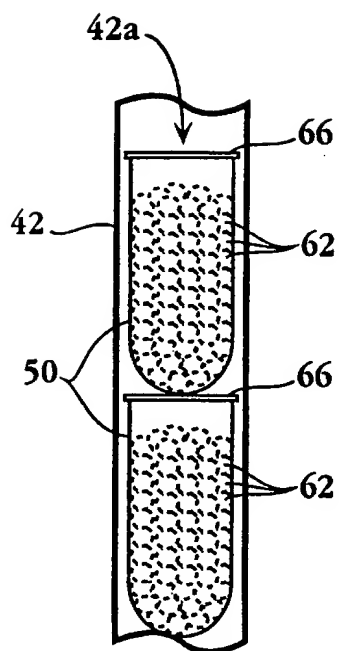


Fig. 4

5/8

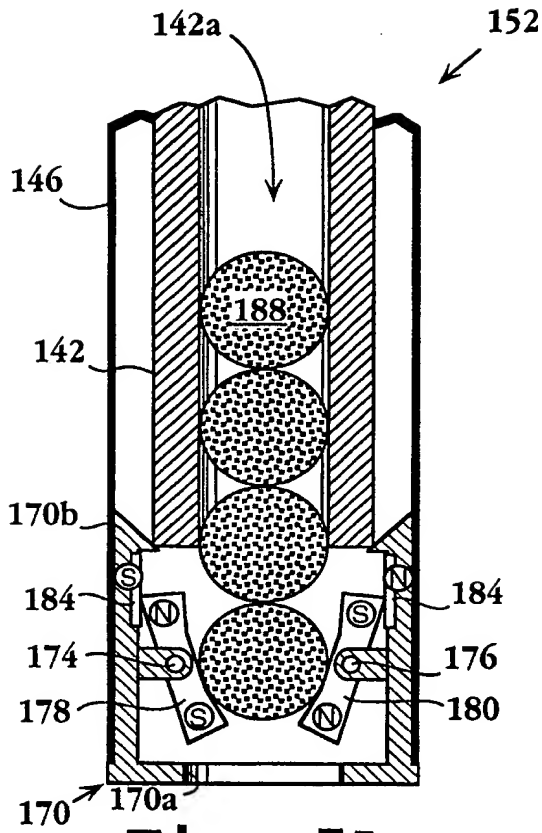


Fig. 5A

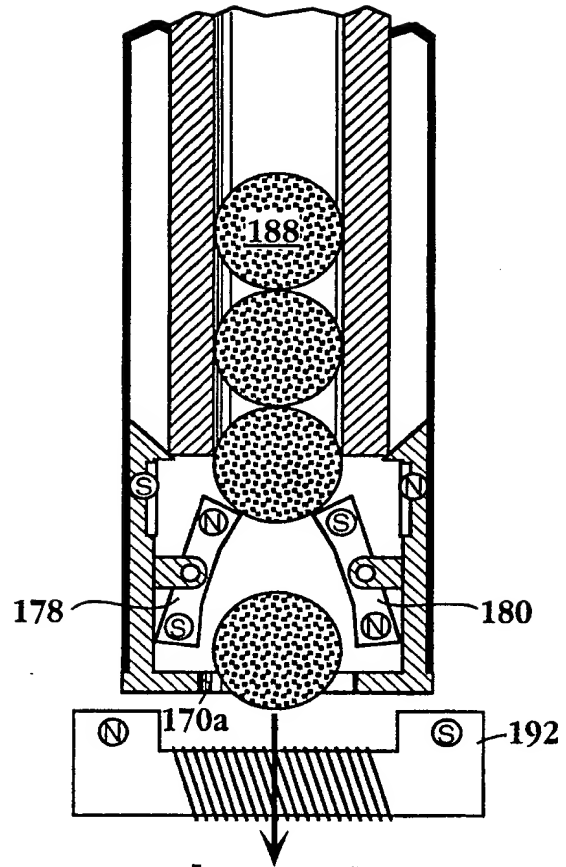


Fig. 6A

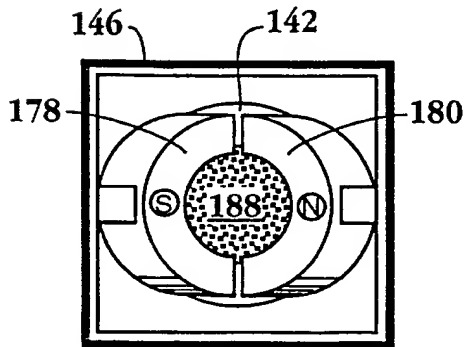


Fig. 5B

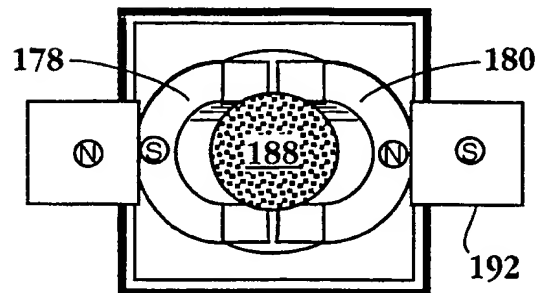


Fig. 6B

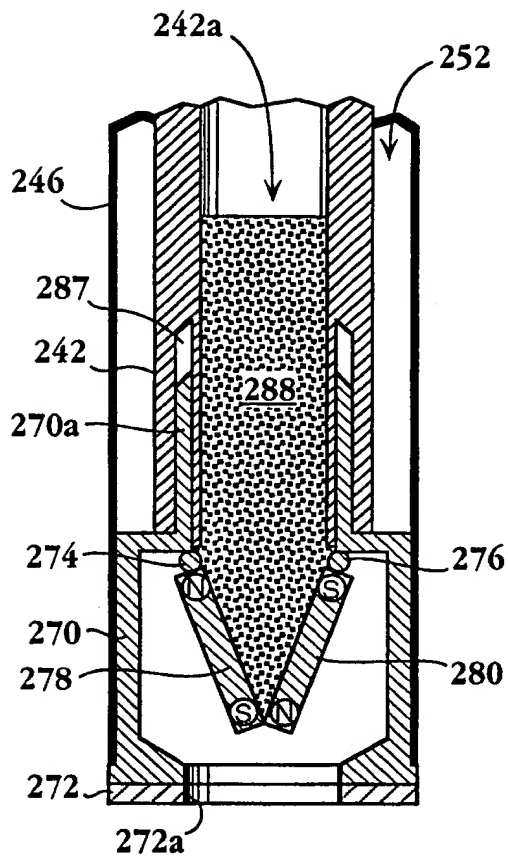


Fig. 7A

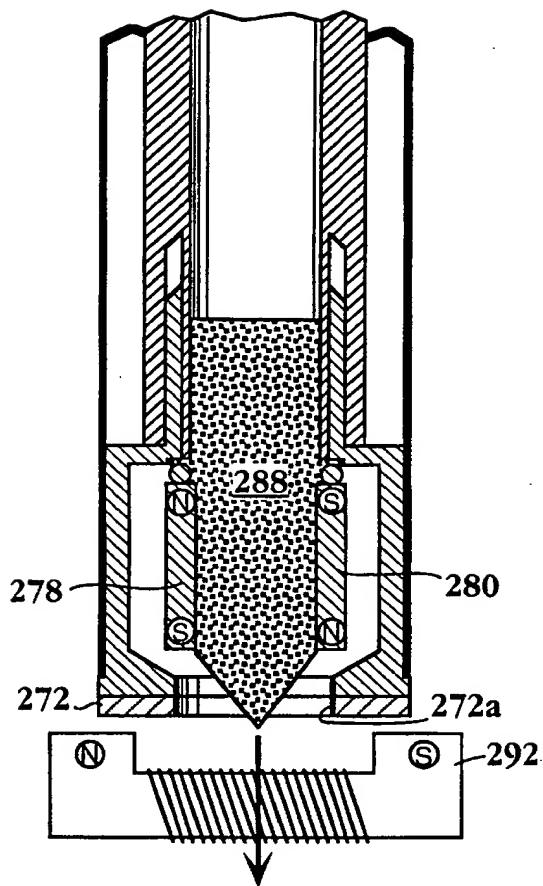


Fig. 8A

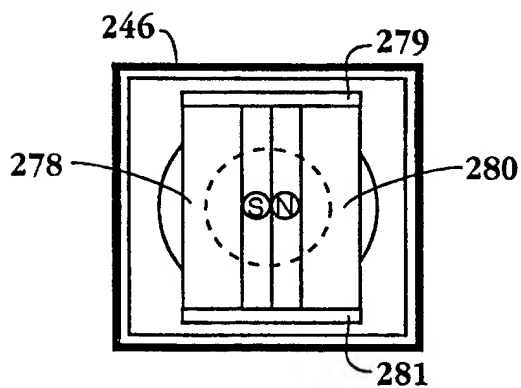


Fig. 7B

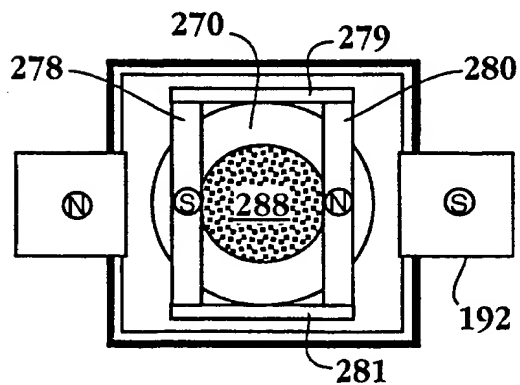
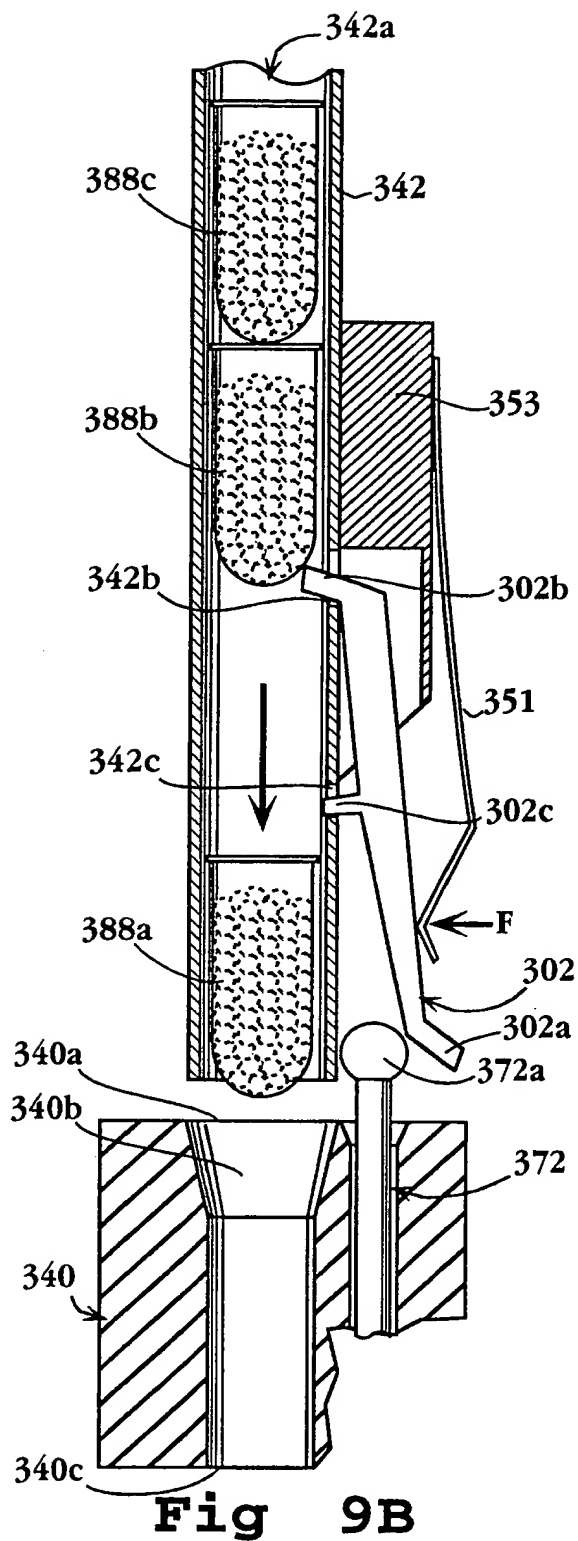
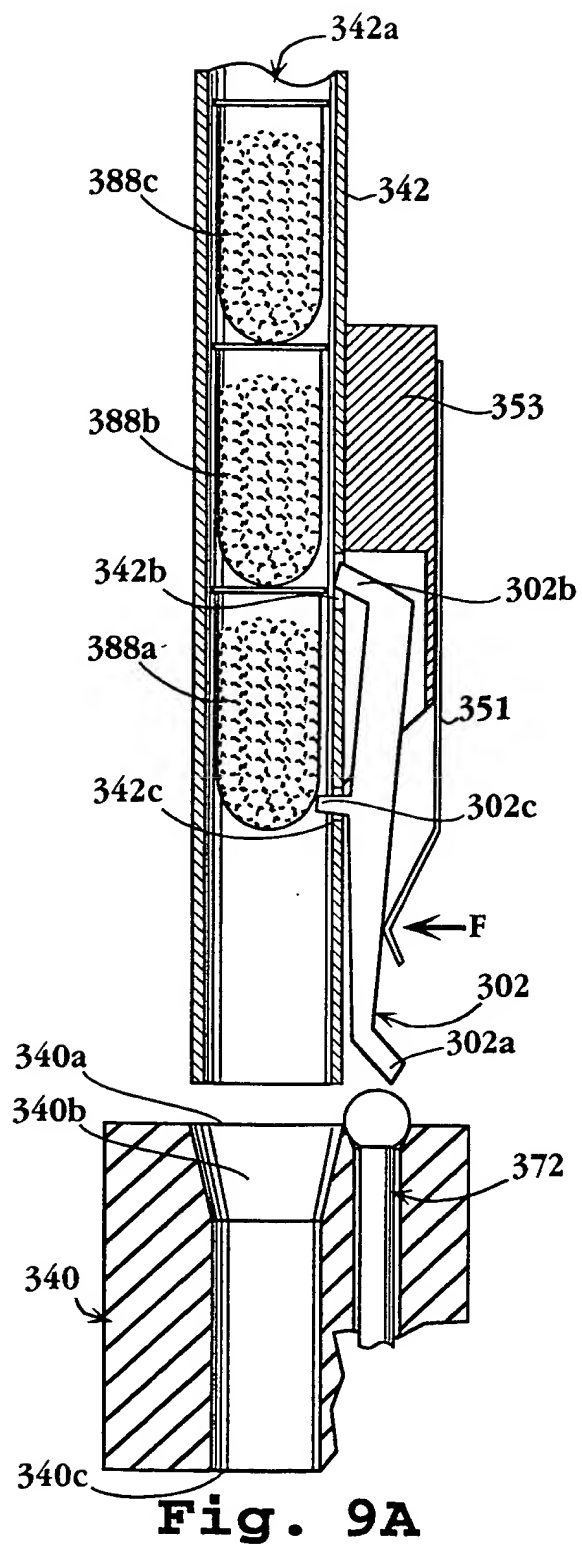


Fig. 8B

7/8



8/8

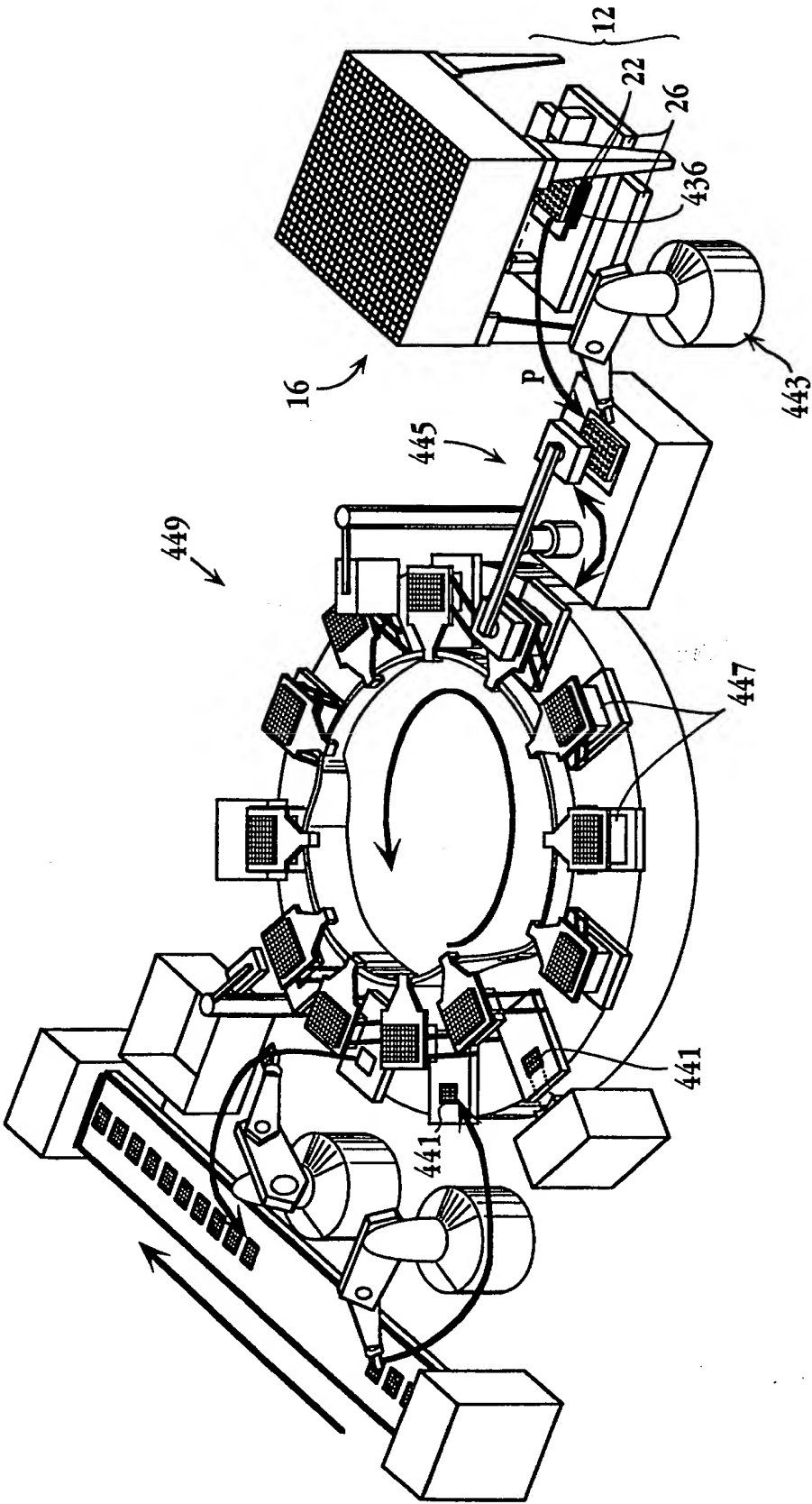


Fig. 10

This Page Blank (uspto)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
24 August 2000 (24.08.2000)

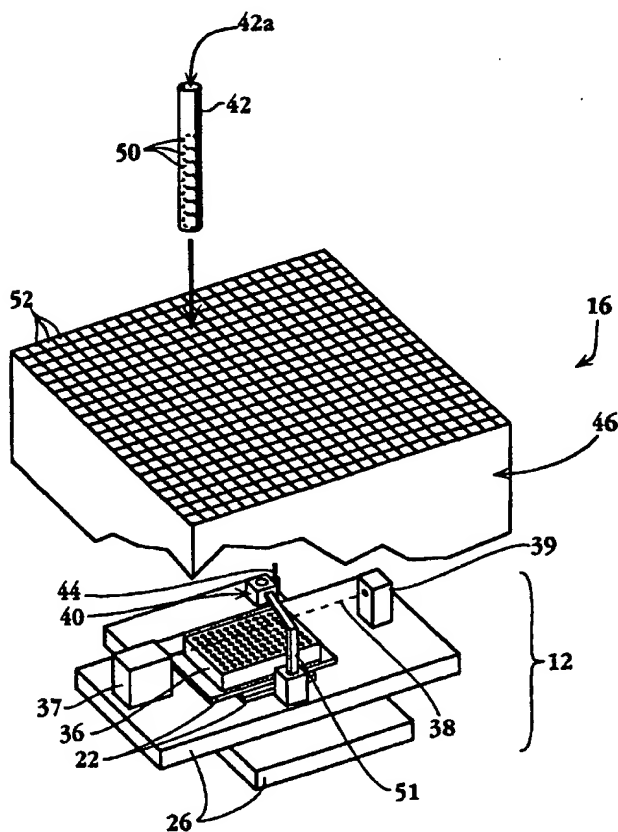
PCT

(10) International Publication Number
WO 00/48735 A3

- (51) International Patent Classification⁷: **B01J 19/00**
- (21) International Application Number: **PCT/US00/03841**
- (22) International Filing Date: **15 February 2000 (15.02.2000)**
- (25) Filing Language: **English**
- (26) Publication Language: **English**
- (30) Priority Data:
09/251,232 16 February 1999 (16.02.1999) **US**
- (63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application:
US 09/251,232 (CIP)
Filed on 16 February 1999 (16.02.1999)
- (71) Applicant (for all designated States except US): **THE PERKIN-ELMER CORPORATION [US/US]; 850 Lincoln Centre Drive, Foster City, CA 94404 (US).**
- (72) Inventors; and
(75) Inventors/Applicants (for US only): **VANN, Charles, S. [US/US]; 1143 Vuelta Olivos, Fremont, CA 94539 (US). COX, David, M. [US/US]; 766 Edgewater Boulevard #302, Foster City, CA 94404 (US).**
- (74) Agents: **GABRIK, Michael, T. et al.; Dehlinger & Associates, P.O. Box 60850, Palo Alto, CA 94306-0850 (US).**
- (81) Designated States (national): **AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.**

[Continued on next page]

(54) Title: **MATRIX STORAGE AND DISPENSING SYSTEM**



(57) Abstract: The present invention provides a system and process providing variable access to, as well as quick and accurate dispensing of, numerous selected reagents from a mass storage arrangement. According to one embodiment, an array of reagent dispensers is supported over a movable platform assembly. The platform assembly aligns a designated receiving receptacle under a selected dispenser of the array so that a respective reagent can be dispensed therein. The dispensers in the matrix have a gate at the lower outlet region. The reagents can be in fluid or solid form, such as beads for solid phase synthesis. Advantageously, the apparatus and process can be carried out under the control of a programmed computer.

WO 00/48735 A3



(84) **Designated States (regional):** ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

(88) **Date of publication of the international search report:**
28 December 2000

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Published:

— *With international search report.*

INTERNATIONAL SEARCH REPORT

In. ational Application No

PCT/US 00/03841

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 B01J19/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 B01J B07C A61J B65D G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, INSPEC, COMPENDEX

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2 099 803 A (GOSHE DR KARABI) 15 December 1982 (1982-12-15) page 2, line 2 - line 35; figures 3-6	25,26
A	US 5 620 853 A (SMETHERS RICK T ET AL) 15 April 1997 (1997-04-15) column 8, line 19 -column 9, line 12; figure 10	1-24, 29-33
A	WO 98 15825 A (IRORI) 16 April 1998 (1998-04-16) page 125, line 14 -page 134, line 4; figures 57,59	1-24, 29-33

-/--

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

15 September 2000

Date of mailing of the international search report

25. 09. 2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Veefkind, V

INTERNATIONAL SEARCH REPORT

Ir. ational Application No

PCT/US 00/03841

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 96 36436 A (IRORI ;DAVID GARY S (US); NOVA MICHAEL P (US); SENYEI ANDREW E (US) 21 November 1996 (1996-11-21) page 75, line 18 -page 76, line 24; figure 7 ---	1-24, 29-33
A	WO 98 40159 A (POKORNY VIT ;TREGA BIOSCIENCES INC (US); KRCHNAK VIKTOR (US); LEBL) 17 September 1998 (1998-09-17) page 29, line 35 -page 30, line 17; figure 19C ---	1-24, 29-33
A	US 5 812 410 A (LION NICHOLAS ET AL) 22 September 1998 (1998-09-22) column 4, line 20 -column 10, line 25; figures 1-14 ---	1-24, 29-33
A	EP 0 803 452 A (FLUIDELEC SA) 29 October 1997 (1997-10-29) the whole document ---	1-24, 29-33
A	US 5 306 510 A (MELTZER WALTER) 26 April 1994 (1994-04-26) abstract; figure 1 ---	1-24, 29-33
A	WO 98 38122 A (SARNOFF CORP) 3 September 1998 (1998-09-03) the whole document ---	25-28
A	EP 0 475 533 A (LAUER LABS B V) 18 March 1992 (1992-03-18) column 11, line 45 -column 12, line 6; figures 1,5 ---	25,26
A	US 4 991 628 A (MAEKELAE KEIJO ET AL) 12 February 1991 (1991-02-12) the whole document -----	25,26

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 00/03841

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☒ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☒ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-24,29-33

Independent claim 1 and dependent claims 2-24, relating to the sytem for storing and dispensing a plurality of reagents. Independent claims 29 and dependent claims 30-33, relating to a method for loading a plurality of receptacles with reagents.

2. Claims: 25-28

Independent claim 25 and dependent claims 26-28, relating to a reagent dispenser assembly comprising a container and a gate mechanism including magnets.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/03841

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB 2099803 A	15-12-1982	NONE	
US 5620853 A	15-04-1997	US 5382512 A AT 191566 T CA 2170249 A DE 69423896 D EP 0715719 A JP 9502519 T WO 9506253 A	17-01-1995 15-04-2000 02-03-1995 11-05-2000 12-06-1996 11-03-1997 02-03-1995
WO 9815825 A	16-04-1998	AU 4976797 A EP 0931259 A AU 6247998 A EP 0959985 A WO 9831732 A	05-05-1998 28-07-1999 07-08-1998 01-12-1999 23-07-1998
WO 9636436 A	21-11-1996	US 5741462 A US 5925562 A US 5751629 A US 5874214 A US 6025129 A AU 707444 B AU 5918596 A AU 7257396 A CA 2216645 A CN 1181720 A EP 0822861 A EP 0853497 A JP 11511238 T WO 9712680 A US 6100026 A US 5961923 A US 6017496 A	21-04-1998 20-07-1999 12-05-1998 23-02-1999 15-02-2000 08-07-1999 29-11-1996 28-04-1997 21-11-1996 13-05-1998 11-02-1998 22-07-1998 28-09-1999 10-04-1997 08-08-2000 05-10-1999 25-01-2000
WO 9840159 A	17-09-1998	US 6045755 A AU 6695098 A	04-04-2000 29-09-1998
US 5812410 A	22-09-1998	US 5838575 A AU 1333097 A WO 9722059 A	17-11-1998 03-07-1997 19-06-1997
EP 0803452 A	29-10-1997	FR 2747900 A	31-10-1997
US 5306510 A	26-04-1994	US 5055263 A	08-10-1991
WO 9838122 A	03-09-1998	US 6047854 A	11-04-2000
EP 0475533 A	18-03-1992	DE 69124556 D DE 69124556 T US 5217590 A	20-03-1997 11-09-1997 08-06-1993
US 4991628 A	12-02-1991	FI 881778 A DE 3910086 A FR 2630180 A GB 2221014 A,B IT 1232482 B JP 1303388 A JP 2014770 C	16-10-1989 23-11-1989 20-10-1989 24-01-1990 17-02-1992 07-12-1989 02-02-1996

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/03841

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4991628 A		JP 7043048 B	15-05-1995